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(54) **Process for preparation of 5-substituted tetrazoles**

Verfahren zur Herstellung von 5-substituierten Tetrazolen

Procédé de préparation de tétrazoles 5-substitués

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Description

[0001] The present invention relates to a process for preparing 5-substituted tetrazoles from nitriles with ease and safety at low costs, said tetrazoles being useful in the fields of pharmaceutical compositions, agricultural compositions, foaming agents, automotive inflators, etc.

[0002] The present invention also concerns with a process for preparing tetrazole derivatives which can be used as an intermediate in the preparation of conventional biphenyl tetrazole derivatives having a high antagonistic activity against angiotensin II.

[0003] Various researches have been conducted on processes for preparing tetrazoles from nitriles and azides, and many processes are known.

[0004] For example, in the synthesis of 5-phenyl-1H-tetrazole, dimethylformamide (DMF) is used as a solvent and NH_4Cl as a catalyst (J. Am. Chem. Soc., 1958, 80, 3908). If NH_4Cl is used as a catalyst in this process, it reacts with NaN_3 to give explosive NH_4N_3 as a sublimate attached to a condenser tube. Thus, the process involves a great risk in industrial manufacture of tetrazoles. However, if NH_4Cl is not used, a tetrazole is produced in a significantly low yield. Stated more specifically, if a catalyst other than NH_4Cl , e.g. LiCl , is used, the yield would be reduced. Further the process requires a high reaction temperature and a prolonged reaction time as a whole.

[0005] In recent years, a process has been known for preparing a tetrazole using an amine salt as a catalyst in DMF solvent. However, if DMF is used as a solvent in the reaction, the DMF itself would decompose due to an alkali component such as an amine. Further, a step of adding water after the reaction is needed to reduce the solubility of a tetrazole in DMF. In this case, since DMF is a watersoluble solvent, unreacted substances would remain in the aqueous layer, consequently necessitating an additional step for separating and removing the unreacted substances from the tetrazole. Moreover, because DMF remains in the mother liquor after the reaction, this would pose problems in industrial manufacture, such as the recovery of organic solvent and the treatment of waste water. That is, the process is industrially markedly disadvantageous.

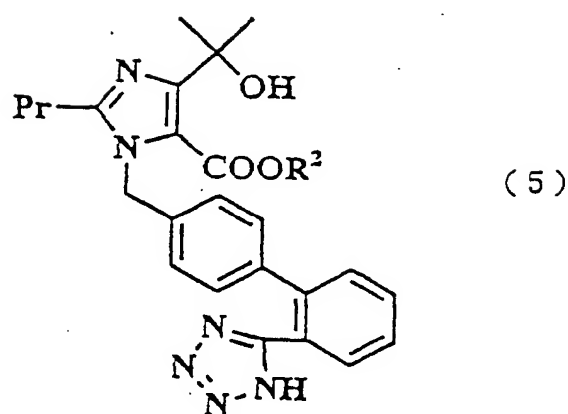
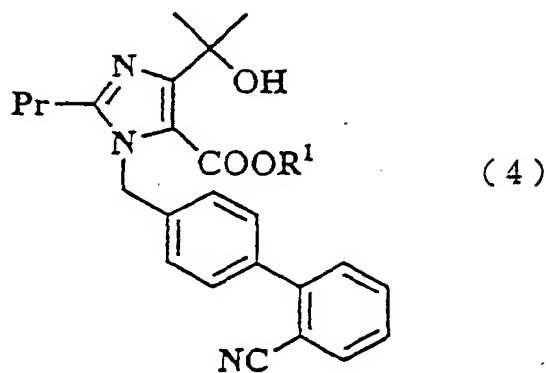
[0006] The reaction would proceed in the presence of a solvent such as aprotic polar solvents other than DMF, and ethers and cellosolves such as tetrahydrofuran, ethylene glycol dimethyl ether, diethylene glycol dimethyl ether, ethylene glycol monomethyl ether and ethylene glycol monoethyl ether. Yet, the reaction is industrially disadvantageous in the reaction rate, reaction ratio and treatment methods.

[0007] A process is known wherein an acid such as acetic acid is used as a catalyst and wherein HN_3 is generated in the presence of a solvent and is reacted with a nitrile (J. Org. Chem., 1957, 22, 1142). However, the process entails a relatively low reaction rate, a high reaction temperature, a prolonged reaction time and a low yield. Furthermore, toxic HN_3 is likely to escape from the reaction system. That is, this process is problematic from the viewpoint of industrial manufacture.

[0008] If DMF is used as a solvent in the preparation of a tetrazole from a nitrile having a complicated structure, the yield of the tetrazole may be pronouncedly reduced because of the reaction of the nitrile with DMF and the decomposition of DMF. In this case, a process is employable which uses an aromatic solvent or polar solvent and an organotin compound, organic silicon compound or the like as a catalyst to increase the organic properties of an azide for the facilitated production of the tetrazole (J. Org. Chem., 1991, 56, 2395). However, the organotin compound used in this process is generally of very high toxicity and the organometallic compound, which is expensive, is required in a large amount compared with the amount of the resulting tetrazole. The obtained tetrazole is often transformed to an oil in the separation from an organometallic compound. In this case, to remove the organometallic compound after conversion of the tetrazole to free form, this necessitates washing or extraction with hexane or like organic solvent, or extraction after treatment of the organometallic compound to enhance its solubility in the solvent for the extraction. Yet in most cases, an organometallic compound can not be completely separated from the tetrazole even by such treatment. In short, if an organometallic compound is used, a tetrazole would be difficult to industrially isolate. Further, from the viewpoint of industrial manufacture, if an organometallic compound is used, various problems are entailed which include requiring time for washing the equipment after the reaction, needing an organic solvent for washing and treating waste water containing the organometallic compound.

[0009] As described above, conventional processes for preparing tetrazoles raise various industrial problems. For example, if NH_4Cl is used, a risk of giving a sublimate is involved. If DMF is used as a solvent, this offers problems such as the degradation of DMF, and the isolation of tetrazole from unreacted nitrile and the treatment of waste water due to the dissolution of DMF in water. Further if an acid is used as a catalyst, this raises problems of a low reaction rate and a low yield. If an organometallic compound is used to increase the yield of a tetrazole, this encounters an industrial problem on the separation of the obtained tetrazole from the organometallic compound.

[0010] Among known processes for preparing 1-(tetrazolylbiphenylmethyl)imidazole derivative of the formula (5) from 1-(cyanobiphenylmethyl)imidazole derivative of the formula (4), a process is known wherein a reaction is made in an aromatic hydrocarbon solvent or a polar solvent in the presence of an organotin azide (Japanese Unexamined Patent Publication No.53,489/1995).



[0011] EP-0264008-A discloses a process for producing 5-methyltetrazole which comprises reacting acetonitrile with an alkali metal azide or ammonium azide in the presence of triethylamine.

[0012] JP-A-93108734 discloses a process for producing a 5-(1,1'-biphenyl)-1H-tetrazole by reacting a 1,1'-biphenyl-2-carbonitrile with an alkali metal azide in the presence of an alkylamine salt.

[0013] In EP-A-0578125 a tri-higher alkyltin azide, $(R)_3SnN_3$, is used to produce tetrazolylbenzene compounds from cyanobenzene compounds.

[0014] US-A-2977372 relates to the production of 5-substituted tetrazoles in a suitable solvent by reaction of an azide salt with an organic nitrile, preferably ammonium azide and dimethyl formamide.

[0015] Methoden der Organischen Chemie, Houben-Weyl, vol E8d (1994) pages 705-8 describes methods for the formation of 5-substituted tetrazoles using nitriles and azides, primarily sodium azide, in several organic solvents.

[0016] DE-A-3829431 discloses particular sulfonamides and several methods for their production. These include the reaction of a substituted nitrile with an azide in the presence of ammonium chloride or alkylammonium chloride in an organic solvent.

[0017] Japanese Unexamined Patent Publication No.53,489/1995 discloses that tributyltin azide synthesized from tributyltin chloride and sodium azide is reacted with 1-(cyanobiphenylmethyl)imidazole derivative of the formula (4) in a toluene solvent, or tributyltin chloride and sodium azide are reacted with 1-(cyanobiphenylmethyl)-imidazole derivative of the formula (4) in a toluene solvent after which the reaction mixture is hydrolyzed in a diluted aqueous solution of sodium hydroxide, followed by neutralization of the aqueous layer with an acid, giving 1-(tetrazolylbiphenylmethyl)imidazole derivative of the formula (5) in a yield of 54%.

[0018] However, the process presents the following problems.

(1) It is difficult to produce pure 1-(tetrazolylbiphenylmethyl)imidazole derivative because of difficulty in completely separating a tributyltin compound from 1-(tetrazolylbiphenylmethyl)imidazole derivative.

(2) The reaction product must be fully purified to remove the tributyltin compound since 1-(tetrazolylbiphenylmethyl)imidazole derivative containing even a small amount of tin can not be used as a pharmaceutical product.

(3) Generally an organotin compound is of high toxicity and calls for care in handling.

(4) Although expensive, an organotin compound is required in a large amount compared with the amount of 1-(tetrazolylbiphenylmethyl)imidazole derivative produced.

(5) For industrial manufacture, a prolonged time is taken for washing to remove the organotin compound and an organic solvent is needed for washing. In addition, a problem is posed for treating waste water containing the organotin compound.

[0019] As stated hereinbefore, conventional processes using an organotin azide pose various problems.

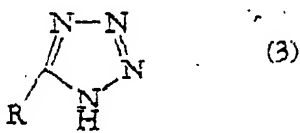
[0020] An object of the present invention is to provide a process capable of preparing a 5-substituted tetrazole in a high yield with ease and safety using inexpensive raw materials while the reaction is easily controlled to inhibit a side reaction.

[0021] It is another object of the invention to provide a process capable of preparing 1-(tetrazolylbiphenylmethyl)imidazole derivative of the formula (5) in a high yield with ease and safety using inexpensive raw materials without use of an organotin compound while the reaction is easily controlled to inhibit a side reaction.

[0022] The inventors of this invention carried out extensive researches on industrially advantageous processes to achieve the foregoing objects and found that an inorganic azide salt is reacted with an amine salt in the presence of an aromatic hydrocarbon as a solvent to give a hydrogen azide salt of amine which is dissolved in the aromatic hydrocarbon. Further findings were that when a hydrogen azide salt of amine is reacted with a nitrite, a tetrazole is produced with high purity in a high yield by a simple procedure and that the solvent can be re-used and the waste water can be easily treated. Based on these novel findings, the present invention was completed.

[0023] Since an inorganic azide salt is scarcely dissolved in an aromatic hydrocarbon in the synthesis of a tetrazole from an inorganic azide salt and a nitrile, a reaction has been considered not to occur in an aromatic hydrocarbon solvent. Consequently, only reactions using a polar solvent have been known.

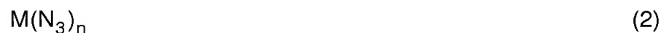
[0024] According to the present invention, there is provided a process for producing a 5-substituted tetrazole of formula (3) :



wherein R is an aliphatic group, an alicyclic group, an aromatic group, a heterocyclic group or a heterocyclic aliphatic group, each group being unsubstituted or substituted by halogen, hydroxy, cyano, nitro, C₁-C₆ alkoxy or amino, or R is C₆-C₁₂ cycloalkyl or branched or unbranched C₁-C₆ alkyl, each of which is unsubstituted or substituted by phenyl, biphenyl or naphthyl, wherein the phenyl, biphenyl and naphthyl groups may be unsubstituted or substituted by halogen, hydroxy, cyano, nitro, C₁-C₆ alkoxy or amino, which process comprises reacting, in an aromatic hydrocarbon solvent in the presence of 1 to 5 moles of an amine salt, 1 mole of a nitrile of formula (1) :



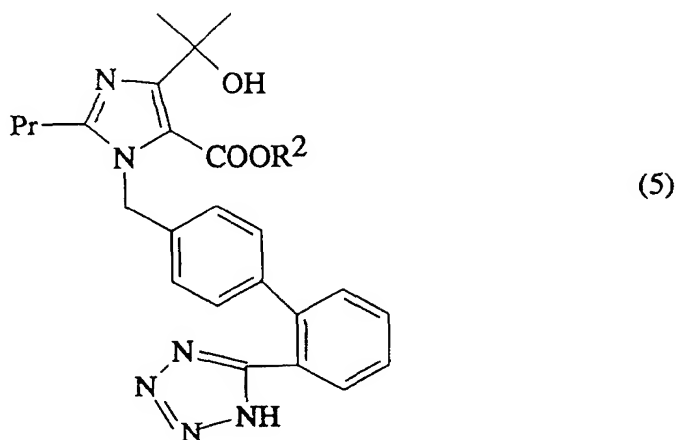
wherein R is as defined above, and 1 to 5 moles of an inorganic azide salt of formula (2):



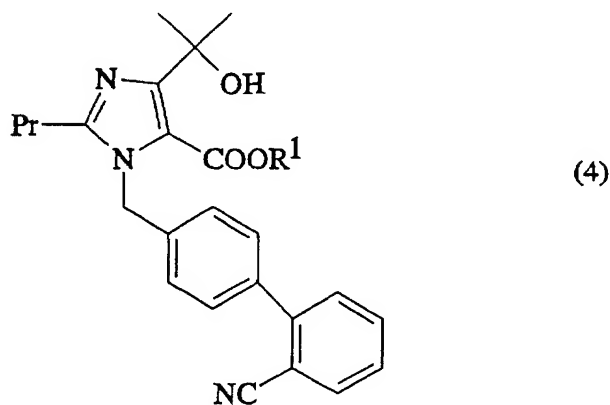
wherein M is an alkali metal or an alkaline earth metal, and n is 1 or 2.

[0025] The present inventors made further investigations on industrially advantageous processes to achieve the foregoing other object and found that when an inorganic azide salt is reacted with an amine salt in an aromatic hydrocarbon as a solvent, a hydrogen azide salt of amine is produced and is dissolved in the aromatic hydrocarbon solvent. More findings were that when the hydrogen azide salt of amine is reacted with 1-(cyanobiphenylmethyl)imidazole derivative of the formula (4), optionally followed by hydrolysis of the obtained compound, 1-(tetrazolylbiphenylmethyl)imidazole derivative of the formula (5) can be produced with high purity in a high yield by a simple procedure, the solvent and the unreacted substances can be re-used, and the waste water can be easily treated. The present invention was completed based on these novel findings.

[0026] According to the present invention, there is provided a process for producing a 1-(tetrazolylbiphenylmethyl)imidazole derivative of formula (5) :



wherein R² is hydrogen or a C₁-C₄ alkyl group, which process comprises reacting, in an aromatic hydrocarbon solvent in the presence of an amine salt, a 1-(cyanobiphenylmethyl)imidazole derivative of formula (4):



wherein R¹ is hydrogen or a C₁-C₄ alkyl group, and an inorganic azide salt of formula (2):



wherein M is an alkali metal or an alkaline earth metal, and n is 1 or 2, and, optionally, hydrolysing the obtained compound.

[0027] Nitriles used as one of the starting materials in the invention include a wide range of nitriles having simple to complicated structures. When in formula (1), R is substituted by a cyano group, the nitrile of the formula (1) is a polycyano compound.

[0028] Examples of an aliphatic group include branched or unbranched C₁-C₃₀ alkyl, typically C₁-C₂₀ alkyl, for instance C₁-C₁₈ alkyl. The said alkyl group may be unsubstituted or substituted by a substituent such as halogen, hydroxy, cyano, nitro, C₁-C₆ alkoxy or amino. As used herein "amino" means a group -NR³R⁴ wherein R³ and R⁴, which are the same or different, are each hydrogen or C₁-C₆ alkyl.

[0029] Examples of an alicyclic group include a C₆-C₁₂ cycloalkyl group such as cyclopentyl, cyclohexyl or cycloheptyl. Examples of an aromatic group include phenyl, biphenyl and naphthyl, each of which may be unsubstituted or

substituted by a substituent such as halogen, hydroxy, cyano, nitro, C₁-C₆ alkoxy or amino.

[0030] Examples of an aromatic aliphatic group include branched or unbranched C₁-C₆ alkyl substituted by phenyl, biphenyl or naphthyl, wherein the phenyl, biphenyl and naphthyl groups may be unsubstituted or substituted by a substituent such as halogen, hydroxy, cyano, nitro, C₁-C₆ alkoxy or amino.

[0031] Examples of an aromatic alicyclic group include C₆-C₁₂ cycloalkyl substituted by an aromatic group as defined above.

[0032] Examples of a heterocyclic group include unsubstituted and substituted, saturated and unsaturated 5- and 6-membered heterocyclic rings containing 1 or more heteroatoms selected from O, S and N, such as furan, thiophene, piperidine, pyrrole, pyrazole, indole and pyridine. The heterocyclic groups may be substituted by substituents such as halogen, hydroxy, cyano, nitro, C₁-C₆ alkoxy or amino.

[0033] Examples of a heterocyclic aliphatic group include branched or unbranched C₁-C₆ alkyl substituted by a heterocyclic group as defined above.

[0034] Specific examples of nitriles which can be used in the invention are acetonitrile, propionitrile, butyronitrile, valeronitrile, hexanenitrile, heptyl cyanide, octyl cyanide, undecanenitrile, dodecanenitrile, tridecanenitrile, pentadecanenitrile, stearonitrile, chloroacetonitrile, bromoacetonitrile, chloropropionitrile, bromopropionitrile, hydroxypropionitrile, methoxyacetonitrile, dimethylaminopropionitrile, dimethylcyanamide, diethylcyanamide, dimethylaminoacetonitrile, cyanoacetamide, cyanoacetic acid, ethyl cyanoformate and like aliphatic nitriles; cyclopropanecarbonitrile, cyclohexanecarbonitrile, cycloheptanecarbonitrile and like alicyclic nitriles; benzonitrile, tolunitrile, cyanophenol, aminobenzonitrile, chlorobenzonitrile, bromobenzonitrile, cyanobenzoic acid, nitrobenzonitrile, anisonitrile, bromotolunitrile, methylcyanobenzoate, m-methoxybenzonitrile, aminotolunitrile, acetylbenzonitrile, naphthonitrile, biphenylcarbonitrile and like aromatic nitriles; phenylacetonitrile, phenylpropionitrile, phenylbutyronitrile, α-methylphenylacetonitrile, methylphenylacetonitrile, diphenylacetonitrile, naphthylacetonitrile, biphenylacetonitrile, aminophenylacetonitrile, nitrophenylacetonitrile, chlorobenzyl cyanide and like aromatic aliphatic nitriles; phenylcyclohexanecarbonitrile, tolylcyclohexanecarbonitrile and like aromatic alicyclic nitriles; cyanofuran, thiophenecarbonitrile, piperidinecarbonitrile, 1,5-dimethyl-2-pyrrolecarbonitrile, pyrazolecarbonitrile, cyanoindole, cyanopyridine and like heterocyclic nitriles; thiopheneacetonitrile, pyridineacetonitrile, N-(2-cyanoethyl)pyrrole, N-methyl-2-pyrroleacetonitrile, indolylacetonitrile and like heterocyclic aliphatic nitriles; cyanobiphenyl derivatives; alkoxycarbonylalkyl cyanide derivatives; and malononitrile, succinonitrile, glutaronitrile, phthalonitrile and like polycyano compounds; to which useful nitriles are not limited. Each nitrile may have a substituent(s). Preferred nitriles are, for example, acetonitrile, benzonitrile, tolunitrile, cyanophenol, aminobenzonitrile, chlorobenzonitrile, phenylacetonitrile, phenylpropionitrile, phenylbutyronitrile, diphenylacetonitrile, cyanopyridine, cyanobiphenyl derivatives, alkoxycarbonylalkyl cyanide derivatives, etc. More preferred nitriles are, for example, acetonitrile, benzonitrile, cyanobiphenyl derivatives, alkoxycarbonylalkyl cyanide derivatives, etc.

[0035] Among 1-(cyanobiphenylmethyl)imidazole derivatives of the formula (4), those wherein R¹ is an alkyl group having 1 to 4 carbon atoms are preferred and those wherein R¹ is a methyl group or an ethyl group are more preferred.

[0036] In the invention, 1-(cyanobiphenylmethyl)-imidazole derivative of the formula (4) can be prepared by any process. An example of the processes is disclosed in Japanese Unexamined Patent Publication No.53,489/1995.

[0037] Examples of useful inorganic azide salts include azides of alkali metals or alkaline earth metals such as sodium, potassium, lithium, calcium, magnesium, etc. Azides of alkali metals are suitable and sodium azide is industrially most suitable.

[0038] The amount of the inorganic azide salt used is 1 to 5 moles, preferably 1 to 3 moles, calculated as hydrogen azide, per mole of the nitrile of the formula (1). Desirably the inorganic azide salt is used in an equimolar amount relative to the amine salt.

[0039] The inorganic azide salt is used in an amount of 1 to 5 moles, preferably 2 to 4 moles, calculated as hydrogen azide, per mole of 1-(cyanobiphenylmethyl)-imidazole derivative of the formula (4). Desirably the inorganic azide salt is used in an equimolar amount relative to the amine salt.

[0040] Amine salts which can be used in the invention are formed from an amine and an acid. Useful amines can be any of primary, secondary and tertiary amines, and aliphatic amines are preferred. Useful amine salts include primary amine salts such as methylamine salt, ethylamine salt, propylamine salt, butylamine salt, amylamine salt, hexylamine salt, cyclohexylamine salt, heptylamine salt, octylamine salt, allylamine salt, benzylamine salt, α-phenylethylamine salt, β-phenylethylamine salt, etc.; secondary amine salts such as dimethylamine salt, diethylamine salt, dipropylamine salt, dibutylamine salt, diamylamine salt, dihexylamine salt, dicyclohexylamine salt, diallylamine salt, morpholine salt, piperidine salt, hexamethyleneimine salt, etc.; and tertiary amine salts such as trimethylamine salt, triethylamine salt, tripropylamine salt, tributylamine salt, triamylamine salt, trihexylamine salt, triallylamine salt, pyridine salt, triethanolamine salt, N-methylmorpholine salt, N,N-dimethylcyclohexylamine salt, N,N-dimethylaniline salt, N,N,N',N'-tetramethylethylenediamine salt, 4-dimethylaminopyridine salt, etc. These amine salts can be used either alone or in combination.

[0041] Acids to be used for forming such salts are those which, together with an amine, are capable of producing a

salt. Useful acids include, for example, inorganic acids such as hydrochloric acid, hydrogen bromide, sulfuric acid, nitric acid, phosphoric acid, boric acid, hydrogen azide, chloric acid, carboxylic acid, hydrogen sulfide and the like; and organic acids such as formic acid, acetic acid, trifluoroacetic acid, propionic acid, oxalic acid, methanesulfonic acid, benzenesulfonic acid, toluenesulfonic acid and the like. Preferred acids are, for example, hydrochloric acid, hydrogen bromide, sulfuric acid, hydrogen azide, acetic acid, trifluoroacetic acid, etc.

[0042] Preferred amine salts include, for example, triethylamine hydrochloride.

[0043] Useful amine salts include commercially available products and salts synthesized by the reaction of an amine with an acid.

[0044] The amount of the amine salt used is sufficient if it is the minimum amount required for the progress of the reaction to progress. The amount is 1 to 5 moles, preferably 1 to 3 moles, calculated as amine per mole of the nitrile of the formula (1). The molar ratio of the amine salt affects the reaction rate and yield. It is especially preferred to use the amine salt in an equimolar amount relative to the inorganic azide salt.

[0045] With respect to 1-(cyanobiphenylmethyl)imidazole derivative of the formula (4), the amine salt is used in an amount of 1 to 5 moles, preferably 2 to 4 moles, calculated as amine per mole of the said derivative. The molar ratio of the amine salt affects the reaction rate and yield. It is especially preferred to use the amine salt in an equimolar amount relative to the inorganic azide salt.

[0046] Solvents preferred in the reaction include aromatic hydrocarbons inert to the reaction. Aromatic hydrocarbons which are industrially suitable and appropriate for the reaction are, for example, benzene, toluene, xylene, mesitylene, ethylbenzene, chlorobenzene, nitrobenzene, cumene, chlorotoluene, etc. among which toluene and xylene are preferred. A mixture of at least two of these aromatic hydrocarbons can be used in the reaction. The amount of the aromatic hydrocarbon used may be a minimum amount which allows the reaction to proceed. The amount is usually 1 to 100 ml, preferably 2 to 50 ml, per gram of the nitrile of the formula (1) or gram of 1-(cyanobiphenylmethyl)imidazole derivative of the formula (4). The solvent may contain about 5% or less of water.

[0047] The reaction temperature is not specifically limited, but can be selected from a wide range of 20 to 150°C, preferably 70 to 140°C although it depends on the combination of starting materials, i.e. nitrile, amine salt and solvent. The reaction time is not critical but usually in the range of 1 to 120 hours, preferably 3 to 50 hours.

[0048] The process of the present invention proceeds at a high rate, giving only a small amount of by-products. That is, the reaction ensures a high efficiency and can produce the desired product in a higher yield than conventional techniques.

[0049] When water is added after the reaction, the 5-substituted tetrazole is present as an amine salt in the aqueous layer. The aromatic hydrocarbon solvent layer can be removed by fractionation. When the reaction is terminated leaving an unreacted nitrile portion dissolved in the aromatic hydrocarbon, that portion can be removed and recovered together with the aromatic hydrocarbon solvent by fractionation. The aromatic hydrocarbon solvent can be easily recovered by distillation and can be re-used. When recovered merely by fractionation, the aromatic hydrocarbon solvent even containing a small amount of water and the unreacted nitrile fraction can be re-used by itself in the reaction.

[0050] After fractionation, an acid is added to the aqueous layer containing the amine salt of the 5-substituted tetrazole to give the 5-substituted tetrazole in free form. In this case, the amine forms a salt, along with the added acid, and the salt is dissolved in water. If the obtained 5-substituted tetrazole in free form is soluble in water, an additional step is necessary for separating the 5-substituted tetrazole in free form from the amine salt (e.g. by extracting the 5-substituted tetrazole in free form with a suitable solvent and distilling off the solvent). On the other hand, if insoluble in water, the 5-substituted tetrazole in free form is filtered and isolated as such, giving a substantially pure 5-substituted tetrazole. Consequently one feature of the invention as described above is that the procedure is very easy and a 5-substituted tetrazole can be safely isolated.

[0051] When R in the formula (3) has a hydrolyzable group, the hydrolyzable group may be converted to a different group on hydrolysis in the post-treatment after the reaction.

[0052] Even when a nitrile having a complicated structure is used, the reaction of the invention easily proceeds without the use of an organometallic compound. Because of the absence of an organometallic compound, the post-treatment after the reaction is easy and inexpensive.

[0053] The process of the invention is improved over techniques using an organometallic compound in safety, toxicity, handleability, costs, ease of procedure, reaction time, yield, etc.

[0054] Especially when using, as the raw material, 1-(cyanobiphenylmethyl)imidazole derivative of the formula (4) wherein R¹ is an alkyl group, the ester moiety may be hydrolyzed with a hydroxide of alkali metal or alkaline earth metal and water after the reaction. Examples of useful hydroxides of alkali metals are lithium hydroxide, sodium hydroxide, potassium hydroxide, etc. Among them, sodium hydroxide is preferred. Useful hydroxides of alkaline earth metals are, for example, barium hydroxide and calcium hydroxide among which calcium hydroxide is favorable. The reaction temperature in the hydrolysis is not critical, but preferably 10 to 40°C. The reaction time in the hydrolysis is not critical but preferably 1 to 5 hours.

[0055] After completion of the hydrolysis, the aqueous layer is washed with an organic solvent such as an aromatic

hydrocarbon and the pH is adjusted to within the acid range so that the 1-(tetrazolylbiphenylmethyl)imidazole derivative of formula (5) wherein R² is a hydrogen atom can be precipitated. The obtained 1-(tetrazolylbiphenylmethyl)-imidazole derivative can be washed with an alcohol, when so required.

[0056] After the reaction, unreacted 1-(cyanobiphenylmethyl)imidazole derivative of the formula (4) remains dissolved in the aromatic hydrocarbon solvent. The remaining derivative can be removed and recovered, along with the aromatic hydrocarbon solvent, by fractionation. The aromatic hydrocarbon solvent can be easily recovered by distillation and can be re-used.

[0057] Since an organotin compound is not used in the invention, tin cannot be present in the 1-(tetrazolylbiphenylmethyl)imidazole derivative of formula (5). Accordingly post-treatment and purification are easy and a high-quality 1-(tetrazolylbiphenylmethyl)imidazole derivative can be easily produced in the invention. One feature of the present invention is that the procedure is easy and the 1-(tetrazolylbiphenylmethyl)imidazole derivative of the formula (5) can be safely isolated at low costs.

[0058] Consequently, the process of the present invention is outstanding in safety, toxicity, handleability, costs, ease of procedure, yield, etc. as compared with processes using an organotin compound.

[0059] As described above, this invention is directed to a significantly advantageous process for preparing a 5-substituted tetrazole on an industrial scale.

[0060] Further, the invention provides a significantly advantageous process for preparing a 1-(tetrazolylbiphenylmethyl)imidazole derivative of the formula (5) on an industrial scale.

[0061] According to the present invention, the 5-substituted tetrazole of formula (3) can be easily and safely prepared from inexpensive raw materials in a high yield while the reaction is easily controlled to inhibit side reactions.

[0062] Moreover, in accordance with the present invention, a 1-(tetrazolylbiphenylmethyl)imidazole derivative of the formula (5) can be easily and safely prepared from inexpensive raw materials in a high yield without use of an organotin compound while the reaction is easily controlled to inhibit side reactions.

[0063] The present invention is described below in more detail with reference to the following Examples. The reaction conditions in Examples are tabulated in Tables 1 to 6. The present invention, however, is not limited to the Examples.

Example 1

[0064] Into a 200-ml flask equipped with a reflux condenser, thermometer and stirrer were placed 5.16 g (0.050 mole) of benzonitrile, 4.23 g (0.065 mole) of sodium azide, 8.95 g (0.065 mole) of triethylamine hydrochloride and 52 ml of toluene. The mixture was heated with stirring to a temperature of 95 to 100°C to undergo a reaction for 7 hours. After completion of the reaction, the reaction mixture was cooled to room temperature and 104 ml of water was added for fractionation. To the aqueous layer was added 6.57 g (0.065 mole) of 36 wt% hydrochloric acid, thereby precipitating 5-phenyl-1H-tetrazole. The precipitate was filtered and dried, giving 7.02 g (0.048 mole) of 5-phenyl-1H-tetrazole (yield 96.0% based on benzonitrile).

Example 2

[0065] A reaction was conducted for 7 hours by following the same procedure as in Example 1 with the exception of using 5.86 g (0.050 mole) of metatolunitrile in place of benzonitrile, whereby 7.46 g (0.047 mole) of 5-(3-methylphenyl)-1H-tetrazole was produced (yield 93.2% based on metatolunitrile).

Example 3

[0066] A reaction was conducted for 24 hours by following the same procedure as in Example 1 with the exception of using 5.86 g (0.050 mole) of orthotolunitrile in place of benzonitrile, whereby 5.05 g (0.032 mole) of 5-(2-methylphenyl)-1H-tetrazole was produced (yield 63.1% based on orthotolunitrile).

Example 4

[0067] A reaction was conducted for 7 hours by following the same procedure as in Example 1 with the exception of using 5.86 g (0.050 mole) of paratolunitrile in place of benzonitrile, whereby 7.73 g (0.048 mole) of 5-(4-methylphenyl)-1H-tetrazole was produced (yield 96.5% based on paratolunitrile).

Example 5

[0068] Into the same device as used in Example 1 were placed 5.96 g (0.050 mole) of orthocyanophenol, 9.76 g (0.150 mole) of sodium azide, 20.65 g (0.150 mole) of triethylamine hydrochloride and 60 ml of toluene. The mixture

was reacted for 5 hours in the same manner as in Example 1, giving 7.95 g (0.049 mole) of 5-(2-hydroxyphenyl)-1H-tetrazole (yield 98.0% based on orthocyanophenol).

Example 6

[0069] Into the same device as used in Example 1 were placed 1.19 g (0.010 mole) of paracyanophenol, 1.95 g (0.030 mole) of sodium azide, 4.13 g (0.030 mole) of triethylamine hydrochloride and 12 ml of toluene. The mixture was reacted for 8 hours in the same manner as in Example 1, giving 1.34 g (0.0083 mole) of 5-(4-hydroxyphenyl)-1H-tetrazole (yield 82.7% based on paracyanophenol).

Example 7

[0070] A reaction was conducted for 1 hour by following the same procedure as in Example 1 with the exception of using 5.86 g (0.050 mole) of phenylacetonitrile in place of benzonitrile and 59 ml of toluene, whereby 5.66 g (0.035 mole) of 5-benzyl-1H-tetrazole was produced (yield 70.7% based on phenylacetonitrile).

Example 8

[0071] A reaction was conducted for 5 hours by following the same procedure as in Example 1 with the exception of using 6.56 g (0.050 mole) of 3-phenylpropionitrile in place of benzonitrile and 66 ml of toluene, whereby 8.16 g (0.047 mole) of 5-phenethyl-1H-tetrazole was produced (yield 93.7% based on 3-phenylpropionitrile).

Example 9

[0072] A reaction was conducted for 7 hours by following the same procedure as in Example 1 with the exception of using 7.26 g (0.050 mole) of 4-phenylbutyronitrile in place of benzonitrile and 73 ml of toluene. After completion of the reaction, the reaction mixture was cooled to room temperature and 74 ml of water was added for fractionation. To the aqueous layer was added 15.22 g (0.030 mole) of 36 wt% hydrochloric acid, and 5-phenylpropyl-1H-tetrazole was extracted with ethyl acetate. The ethyl acetate layer was washed with a 10 wt% aqueous solution of sodium chloride, dried over magnesium sulfate, filtered, and distilled to remove ethyl acetate. This procedure gave 8.03 g (0.043 mole) of 5-phenylpropyl-1H-tetrazole (yield 85.3% based on 4-phenylbutyronitrile).

Example 10

[0073] A reaction was conducted for 23 hours by following the same procedure as in Example 1 with the exception of using 5.91 g (0.050 mole) of 2-aminobenzonitrile in place of benzonitrile, whereby 7.32 g (0.045 mole) of 5-(2-aminophenyl)-1H-tetrazole was produced (yield 90.8% based on 2-aminobenzonitrile).

Example 11

[0074] A reaction was conducted for 28 hours by following the same procedure as in Example 1 with the exception of using 5.91 g (0.050 mole) of 3-aminobenzonitrile in place of benzonitrile, whereby 7.63 g (0.047 mole) of 5-(3-aminophenyl)-1H-tetrazole was produced (yield 94.7% based on 3-aminobenzonitrile).

Example 12

[0075] Into the same device as used in Example 1 were placed 6.58 g (0.065 mole) of triethylamine and 52 ml of toluene. To the mixture were added 3.19 g (0.033 mole) of concentrated sulfuric acid with stirring. Further added were 5.16 g (0.050 mole) of benzonitrile and 4.23 g (0.065 mole) of sodium azide. The mixture was heated to a temperature from 95 to 100°C to undergo a reaction for 7 hours. The same subsequent treatment as in Example 1 was conducted, giving 7.11 g (0.049 mole) of 5-phenyl-1H-tetrazole (yield 97.3% based on benzonitrile).

Example 13

[0076] A reaction was conducted for 8 hours by following the same procedure as in Example 12 with the exception of using 3.91 g (0.065 mole) of glacial acetic acid in place of concentrated sulfuric acid, whereby 6.17 g (0.042 mole) of 5-phenyl-1H-tetrazole was produced (yield 84.4% based on benzonitrile).

Example 14

[0077] Into the same device as used in Example 1 were placed 5.16 g (0.050 mole) of benzonitrile, 4.23 g (0.065 mole) of sodium azide, 7.13 g (0.065 mole) of diethylamine hydrochloride and 52 ml of toluene. The mixture was reacted for 8 hours in the same manner as in Example 1, giving 7.02 g (0.048 mole) of 5-phenyl-1H-tetrazole (yield 96.0% based on benzonitrile).

Example 15

[0078] A reaction was conducted for 8 hours by following the same procedure as in Example 14 with the exception of using 5.30 g (0.065 mole) of ethylamine hydrochloride in place of diethylamine hydrochloride, whereby 4.10 g (0.028 mole) of 5-phenyl-1H-tetrazole was produced (yield 56.1% based on benzonitrile).

Example 16

[0079] Into the same device as used in Example 1 were placed 5.16 g (0.050 mole) of benzonitrile, 4.23 g (0.065 mole) of sodium azide, 8.95 g (0.065 mole) of triethylamine hydrochloride, 52 ml of toluene and 2.6 g of water. The mixture was reacted at a temperature of 85 to 95°C for 8 hours, giving 6.12 g (0.042 mole) of 5-phenyl-1H-tetrazole (yield 83.7% based on benzonitrile).

Example 17

[0080] A reaction was conducted at 76 to 79°C for 8 hours by following the same procedure as in Example 1 with the exception of using 52 ml of benzene in place of toluene, whereby 6.67 g (0.046 mole) of 5-phenyl-1H-tetrazole was produced (yield 91.2% based on benzonitrile).

Example 18

[0081] A reaction was conducted at a temperature of 100 to 107°C for 5 hours by following the same procedure as in Example 1 with the exception of using 52 ml of xylene in place of toluene, whereby 7.22 g (0.049 mole) of 5-phenyl-1H-tetrazole was produced (yield 98.8% based on benzonitrile).

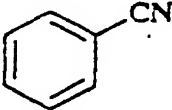
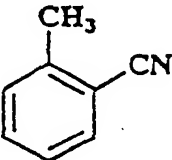
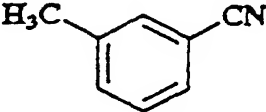
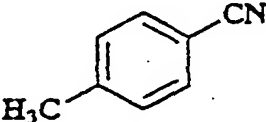

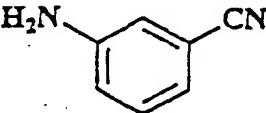
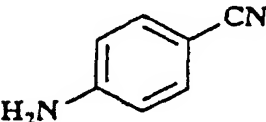
Example 19

[0082] A reaction was conducted at a temperature of 100 to 102°C for 8 hours by following the same procedure as in Example 1 with the exception of using 52 ml of nitrobenzene in place of toluene, whereby 7.19 g (0.049 mole) of 5-phenyl-1H-tetrazole was produced (yield 98.4% based on benzonitrile).

[0083] Tables 1 to 6 show Examples wherein various types of solvents were used in varied amounts, and various types of inorganic azide salts and amine salts were used in varied amounts.

[0084] Table 1 shows Examples wherein reactions were made using various types of nitriles under varied reaction conditions (reaction temperature and reaction time), while using about 10 ml of toluene per gram of a nitrile, and sodium azide and triethylamine hydrochloride respectively in an amount of 1.3 moles per mole of a nitrile (in Example 20, a reaction was performed in the same manner as in Examples 1 to 4, 10 and 11 except that the type of nitriles and reaction conditions were employed as shown in Table 1).

Table 1

| Ex. No. | Nitrile | Reaction Conditions | Yield(%) |
|---------|---|---------------------|----------|
| 1 |  | 95~100°C × 7 hr | 96.0 |
| 3 |  | 95~100°C × 24hr | 63.1 |
| 2 |  | 95~100°C × 7 hr | 93.2 |
| 4 |  | 95~100°C × 7 hr | 96.5 |
| 10 |  | 95~100°C × 23hr | 90.8 |
| 11 |  | 95~100°C × 28hr | 94.7 |
| 20 |  | 95~100°C × 23hr | 92.2 |

[0085] Table 2 shows Examples wherein reactions were made using about 10 ml of toluene per gram of a nitrile, and sodium azide and triethylamine hydrochloride respectively in an amount of 3.0 moles per mole of a nitrile, and employing

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various types of nitriles under varied reaction conditions (in Examples 21 to 25, reactions were conducted in the same manner as in Examples 5 and 6 with the exception of using various types of nitriles and reaction conditions as shown in Table 2).

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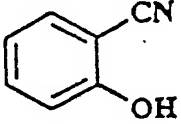
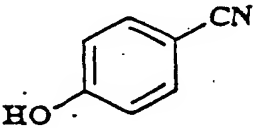
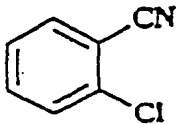
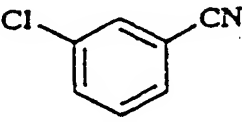
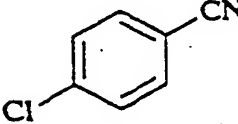
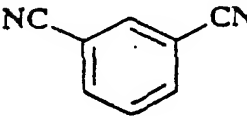
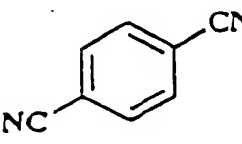
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Table 2

| Ex. No. | Nitrile | Reaction Conditions | Yield(%) |
|---------|---|---------------------|----------|
| 5 |  | 95~100°C × 5 hr | 98.0 |
| 6 |  | 95~100°C × 8 hr | 82.7 |
| 21 |  | 95~100°C × 7 hr | 68.9 |
| 22 |  | 95~100°C × 7 hr | 100 |
| 23 |  | 95~100°C × 7 hr | 99.9 |
| 24 |  | 95~100°C × 7 hr | 96.9 |
| 25 |  | 95~100°C × 7 hr | 100 |

[0086] Table 3 shows Examples wherein reactions were made using various types of nitriles under varied reaction

conditions, while employing about 10 to about 40 ml of toluene per gram of a nitrile, and sodium azide and triethylamine hydrochloride respectively in an amount of 3.0 moles per mole of a nitrile (in Examples 26 to 30, reactions were made in the same manner as in Examples 7 to 9 except that the type of nitriles and reaction conditions were varied as shown in Table 3 and the amount of toluene solvent was varied).

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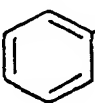

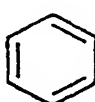
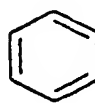
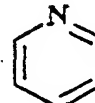
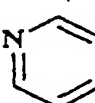
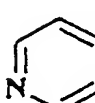
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Table 3

| Ex. No. | Nitrile | Reaction Conditions | Yield(%) |
|---------|---|---|----------|
| 2 6 | CH_3CN | $95\sim 100^\circ\text{C} \times 25\text{hr}$ | 9 0 . 0 |
| 7 |  | $95\sim 100^\circ\text{C} \times 1\text{hr}$ | 7 0 . 7 |
| 8 |  | $95\sim 100^\circ\text{C} \times 5\text{hr}$ | 9 3 . 7 |
| 9 |  | $95\sim 100^\circ\text{C} \times 7\text{hr}$ | 8 5 . 3 |
| 2 7 |  | $95\sim 100^\circ\text{C} \times 7\text{hr}$ | 4 7 . 6 |
| 2 8 |  | $95\sim 100^\circ\text{C} \times 7\text{hr}$ | 1 0 0 |
| 2 9 |  | $95\sim 100^\circ\text{C} \times 7\text{hr}$ | 1 0 0 |
| 3 0 |  | $95\sim 100^\circ\text{C} \times 7\text{hr}$ | 1 0 0 |

[0087] Table 4 shows Examples wherein reactions were made using various types of solvents under varied reaction

conditions, while employing about 10 ml of a solvent per gram of benzonitrile, and sodium azide and triethylamine hydrochloride respectively in an amount of 1.3 moles per mole of benzonitrile. "P5T" is an abbreviation for 5-phenyl-1H-tetrazole (same hereinafter).

Table 4

| Ex. No. | Solvent | Reaction conditions | P5T yield (%) |
|---------|--------------------------|---------------------|---------------|
| 17 | Benzene | 76-79°C X 8 hr | 91.2 |
| 1 | Toluene | 95-100°C X 7 hr | 96.0 |
| 18 | Xylene | 100-107°C X 5 hr | 98.8 |
| 19 | Nitrobenzene | 100-102°C X 8 hr | 98.4 |
| 16 | Toluene + 5% water (w/v) | 85-98°C X 8 hr | 83.7 |

[0088] Table 5 shows Examples wherein reactions were made using different types of catalysts under different reaction conditions, while using about 10 ml of toluene per gram of benzonitrile, and sodium azide and amine salt (catalyst) respectively in an amount of 1.3 moles per mole of benzonitrile.

Table 5

| Ex.No. | Catalyst | Reaction conditions | P5T yield (%) |
|--------|------------------------|---------------------|---------------|
| 1 | Et ₃ N·HCl | 95-100°C X 7 hr | 96.0 |
| 14 | Et ₂ NH·HCl | 95-100°C X 8 hr | 96.0 |
| 15 | EtNH ₂ ·HCl | 95-100°C X 8 hr | 56.1 |

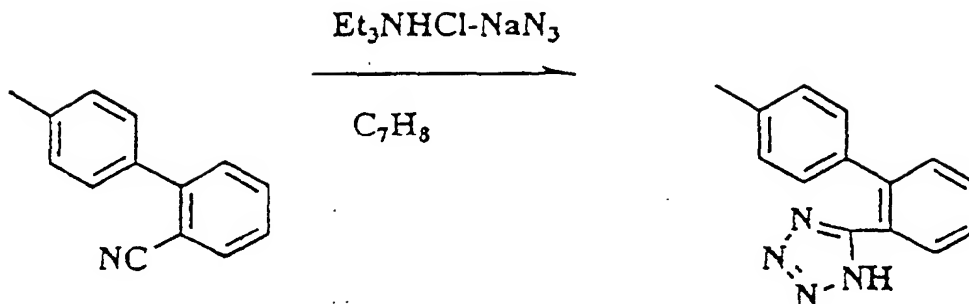
[0089] Table 6 shows Examples wherein reactions were made using various types of acids to form an amine salt under varied reaction conditions, while using about 10 ml of toluene per gram of benzonitrile, and sodium azide and triethylamine respectively in an amount of 1.3 moles per mole of benzonitrile. The molar ratio of the acid is relative to benzonitrile. In Example 1, triethylamine hydrochloride was used, but the acid component is indicated in Table 6.

Table 6

| Ex. No. | Acid(molar ratio) | Reaction conditions | P5T yield (%) |
|---------|-------------------------|---------------------|---------------|
| 1 | Hydrochloric acid (1.3) | 95-100°C X 7 hr | 96.0 |
| 12 | Sulfuric acid (0.66) | 95-100°C X 7 hr | 97.3 |
| 13 | Acetic acid (1.3) | 95-100°C X 8 hr | 84.4 |

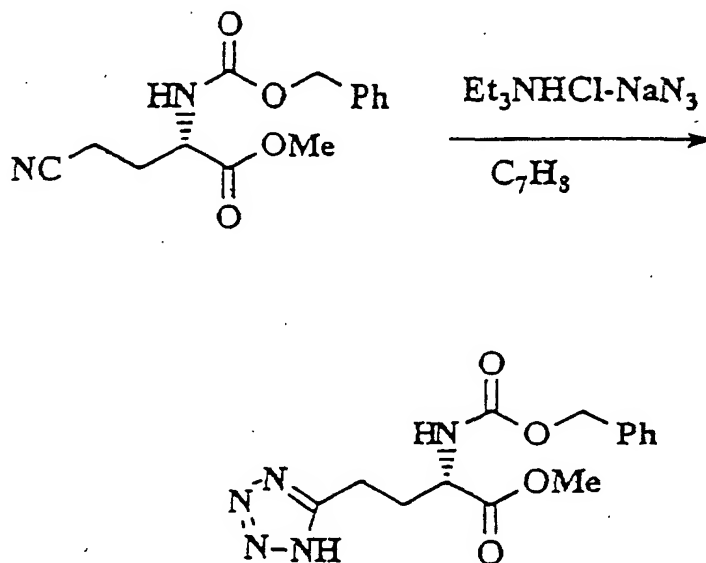
Example 31

[0090] Into the same device as used in Example 1 were placed 9.66 g (0.050 mole) of 4'-methylbiphenyl-2-carbonitrile, 6.50 g (0.100 mole) of sodium azide, 13.77 g (0.100 mole) of triethylamine hydrochloride and 96.6 ml of toluene. The mixture was reacted for 48 hours in the same manner as in Example 1, giving 10.85 g (0.046 mole) of 5-(4'-methylbiphenyl-2-yl)-1H-tetrazole (yield 91.9% based on 4'-methylbiphenyl-2-carbonitrile). The reaction is illustrated below with chemical formulas:



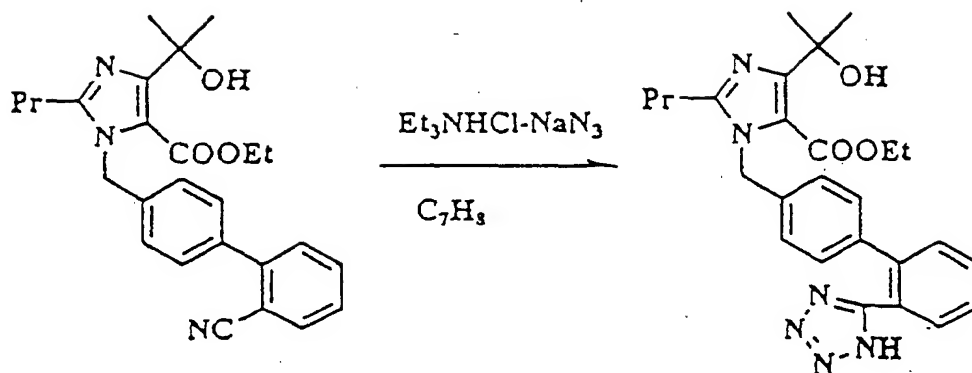
Example 32

[0091] Into the same device as used in Example 1 were placed 77.3 g (0.28 mole) of methyl (2S)-2-benzyloxycarbonylamino-4-cyanobutyrate, 27.3 g (0.42 mole) of sodium azide, 57.7 g (0.42 mole) of triethylamine hydrochloride and 280 ml of toluene. Thereafter, the mixture was heated to a temperature of 82 to 83°C to undergo a reaction for 24 hours. Subsequently the same procedure as in Example 1 was repeated, and crystals were collected by filtration. Further, the crystals were recrystallized from methanol and water, filtered and dried, giving 67.7 g (0.21 mole) of methyl (2S)-2-benzyloxycarbonylamino-4-(tetrazole-5-yl)butyrate (yield 75.8% based on methyl (2S)-2-benzyloxycarbonylamino-4-cyanobutyrate). The reaction is illustrated below with chemical formulas:



Example 33

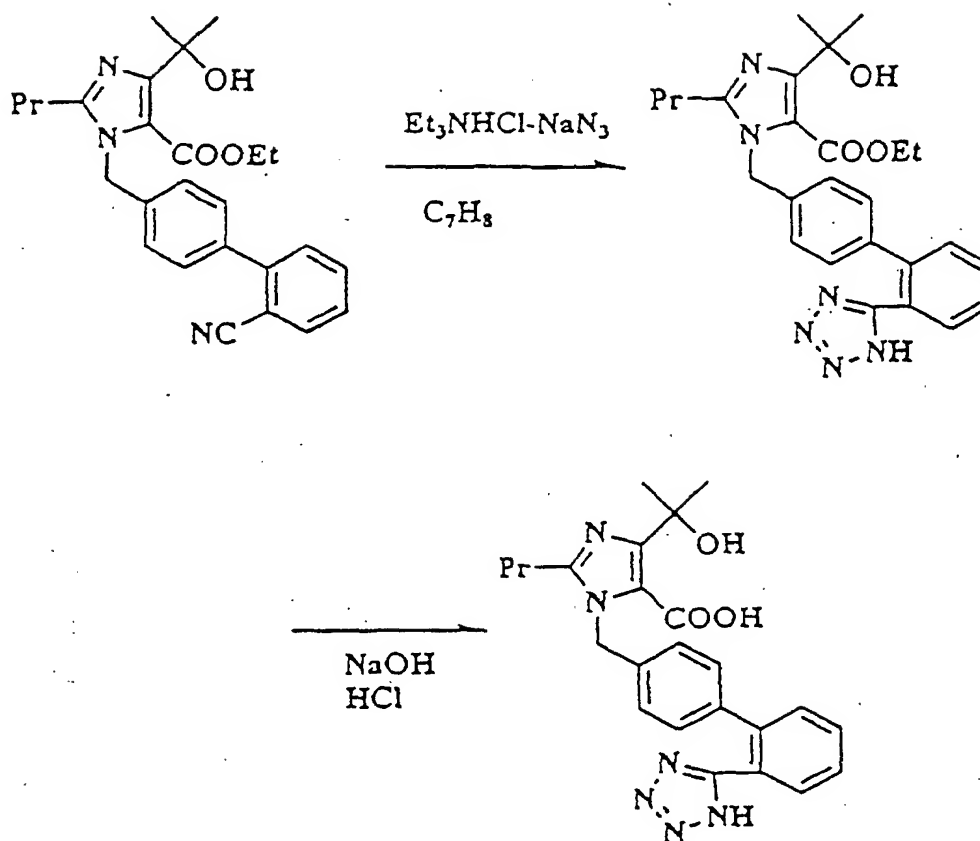
[0092] Into a 500-ml flask equipped with a reflux condenser, thermometer and stirrer were placed 43.5 g (0.10 mole) of ethyl ester of 1-(2'-cyanobiphenyl-4-yl)methyl-4-(1-hydroxy-1-methylethyl)-2-propylimidazole-5-carboxylic acid, 19.5 g (0.30 mole) of sodium azide, 41.3 g (0.30 mole) of triethylamine hydrochloride and 206 ml of toluene. The mixture was heated with stirring to a temperature of 95 to 100°C to undergo a reaction for 24 hours. After completion of the reaction, the reaction mixture was cooled to room temperature. Then, the reaction mixture was analyzed by high-performance liquid chromatography and identified as ethyl ester of 4-(1-hydroxy-1-methylethyl)-2-propyl-1-{4-[2-{tetrazole-5-yl}phenyl]phenyl}methylimidazole-5-carboxylic acid (reaction ratio 87%). The reaction is illustrated below with chemical formulas:



Example 34

[0093] Into a 500-ml flask equipped with a reflux condenser, thermometer and stirrer were placed 43.5 g (0.10 mole) of ethyl ester of 1-(2'-cyanobiphenyl-4-yl)methyl-4-(1-hydroxy-1-methylethyl)-2-propylimidazole-5-carboxylic acid, 19.5 g (0.30 mole) of sodium azide, 41.3 g (0.30 mole) of triethylamine hydrochloride and 206 ml of toluene. The mixture was heated with stirring to a temperature of 95 to 100°C to undergo a reaction for 24 hours. After completion of the reaction, the reaction mixture was cooled to room temperature. Then, 35.0 g (0.42 mole) of a 48 wt% aqueous solution of NaOH and 335 ml of water were added.

[0094] The mixture was fractionated after 2 hours' stirring at room temperature. The aqueous layer was washed and fractionated with the addition of 103 ml of toluene. The aqueous layer was adjusted to a pH of 2 to 3.5 with 36 wt% hydrochloric acid to separate out crystals, which were collected by filtration and dried. The crystals were heated, washed with methanol with stirring, cooled, filtered for collection and dried, giving 32.6 g (0.07 mole) of 4-(1-hydroxy-1-methylethyl)-2-propyl-1-[4-[2-(tetrazole-5-yl)phenyl]phenyl]methylimidazole-5-carboxylic acid (yield 72.4% based on ethyl ester of 1-(2'-cyanobiphenyl-4-yl)methyl-4-(1-hydroxy-1-methylethyl)-2-propylimidazole-5-carboxylic acid). The reaction is illustrated below with chemical formulas.



Example 35

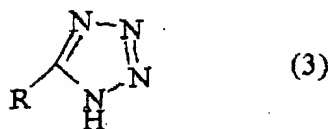
[0095] Into the same device as used in Example 34 were placed 43.5 g (0.10 mole) of ethyl ester of 1-(2'-cyanobiphenyl-4-yl)methyl-4-(1-hydroxy-1-methylethyl)-2-propylimidazole-5-carboxylic acid, 13.1 g (0.20 mole) of sodium azide, 31.9 g (0.23 mole) of triethylamine hydrochloride and 206 ml of toluene. Thereafter, the mixture was reacted and treated in the same manner as in Example 34, giving 27.0 g (0.06 mole) of 4-(1-hydroxy-1-methylethyl)-2-propyl-1-[4-[2-(tetrazole-5-yl)phenyl]phenyl]methyylimidazole-5-carboxylic acid (yield 60.0% based on ethyl ester of 1-(2'-cyanobiphenyl-4-yl)methyl-4-(1-hydroxy-1-methylethyl)-2-propylimidazole-5-carboxylic acid).

Example 36

[0096] Into the same device as used in Example 34 were placed 43.5 g (0.10 mole) of ethyl ester of 1-(2'-cyanobiphenyl-4-yl)methyl-4-(1-hydroxy-1-methylethyl)-2-propylimidazole-5-carboxylic acid, 8.5 g (0.13 mole) of sodium azide, 20.8 g (0.15 mole) of triethylamine hydrochloride and 206 ml of toluene. The mixture was reacted and treated in the same manner as in Example 34, giving 20.3 g (0.045 mole) of 4-(1-hydroxy-1-methylethyl)-2-propyl-1-[4-[2-(tetrazole-5-yl)phenyl]phenyl]methyylimidazole-5-carboxylic acid (yield 45.0% based on ethyl ester of 1-(2'-cyanobiphenyl-4-yl)methyl-4-(1-hydroxy-1-methylethyl)-2-propylimidazole-5-carboxylic acid).

Claims

1. A process for producing a 5-substituted tetrazole of formula (3):



10 wherein R is an aliphatic group, an alicyclic group, an aromatic group, a heterocyclic group or a heterocyclic aliphatic group, each group being unsubstituted or substituted by halogen, hydroxy, cyano, nitro, C₁-C₆ alkoxy or amino, or R is C₆-C₁₂ cycloalkyl or branched or unbranched C₁-C₆ alkyl, -each of which is unsubstituted or substituted by phenyl, biphenyl or naphthyl, wherein the phenyl, biphenyl and naphthyl groups may be unsubstituted or substituted by halogen, hydroxy, cyano, nitro, C₁-C₆ alkoxy or amino, which process comprises reacting, in an aromatic hydrocarbon solvent in the presence of 1 to 5 moles of an amine salt, 1 mole of a nitrite of formula. (1):

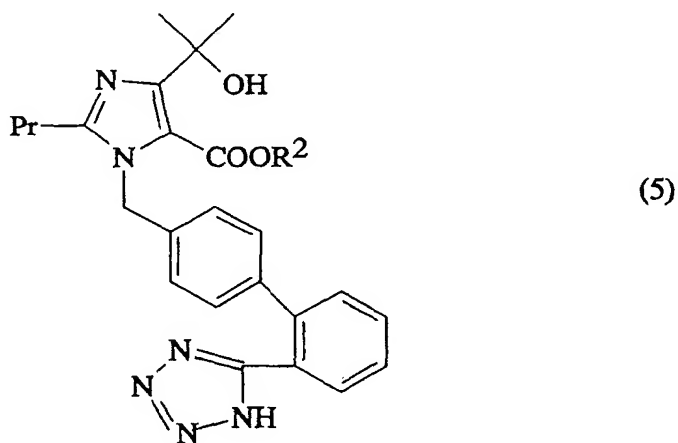


20 wherein R is as defined above, and 1 to 5 moles of an inorganic azide salt of formula (2):

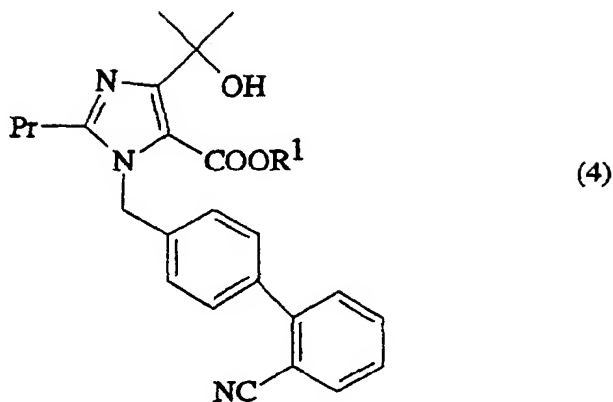


25 wherein M is an alkali metal or an alkaline earth metal, and n is 1 or 2.

2. A process according to claim 1, wherein the inorganic azide salt is sodium azide.
3. A process according to claim 1 or 2, wherein the amine salt is triethylamine hydrochloride.
4. A process according to any one of the preceding claims, wherein the aromatic hydrocarbon solvent is toluene or xylene.
5. A process for producing a 1-(tetrazolylbiphenylmethyl)imidazole derivative of formula (5):



55 wherein R² is hydrogen or a C₁-C₄ alkyl group, which process comprises reacting, in an aromatic hydrocarbon solvent in the presence of 1 to 5 moles of an amine salt, 1 mole of a 1-(cyanobiphenylmethyl)imidazole derivative of formula (4):



wherein R¹ is hydrogen or a C₁-C₄ alkyl group, and 1 to 5 moles of an inorganic azide salt of formula (2) :

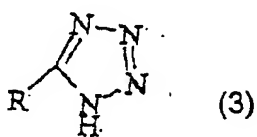


wherein M is an alkali metal or an alkaline earth metal, and n is 1 or 2, and, optionally, hydrolysing the obtained compound.

- 25
6. A process according to claim 5, wherein R¹ in formula (4) is methyl or ethyl and R² in formula (5) is hydrogen.
7. A process according to claim 5 or 6, wherein the inorganic azide salt is sodium azide.
8. A process according to any one of claims 5 to 7 wherein the amine salt is triethylamine hydrochloride.
- 30
9. A process according to any one of claims 5 to 8, wherein the aromatic hydrocarbon solvent is toluene or xylene.

Patentansprüche

- 35
1. Verfahren zur Herstellung eines 5-substituierten Tetrazols der Formel (3):



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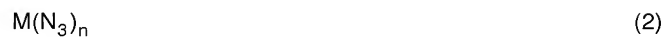
worin R eine aliphatische Gruppe, eine alicyclische Gruppe, eine aromatische Gruppe, eine heterocyclische Gruppe oder eine heterocyclische aliphatische Gruppe ist, wobei jede Gruppe unsubstituiert oder mit Halogen, Hydroxy, Cyano, Nitro, C₁-C₆-Alkoxy oder Amino substituiert ist, oder R C₆-C₁₂-Cycloalkyl oder verzweigtes oder unverzweigtes C₁-C₆-Alkyl ist, die jeweils unsubstituiert oder mit Phenyl, Biphenyl oder Naphthyl substituiert sind, worin die Phenyl-, Biphenyl- und Naphthylgruppen unsubstituiert oder mit Halogen, Hydroxy, Cyano, Nitro, C₁-C₆-Alkoxy oder Amino substituiert sein können, wobei das Verfahren umfaßt die Umsetzung von 1 mol eines Nitrils der Formel

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(1):

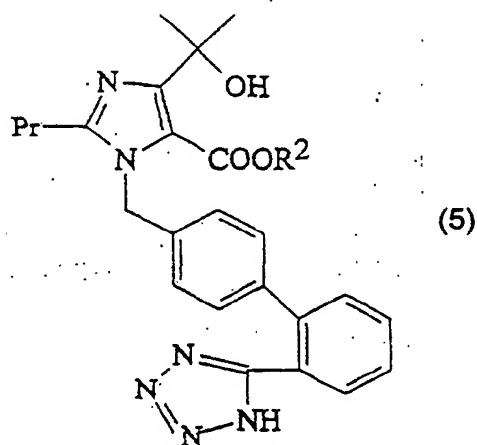


worin R wie vorstehend definiert ist, und 1 bis 5 mol eines anorganischen Azidsalzes der Formel (2):

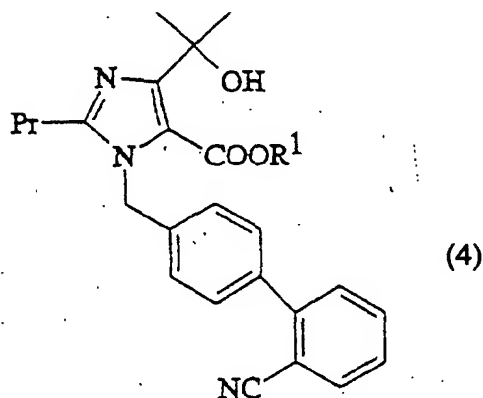


worin M ein Alkalimetall oder ein Erdalkalimetall ist und n 1 oder 2 ist, in einem aromatischen Kohlenwasserstoff-Lösungsmittel in Anwesenheit von 1 bis 5 mol eines Aminsalzes.

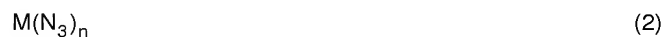
2. Verfahren nach Anspruch 1, worin das anorganische Azidsalz Natriumazid ist.
3. Verfahren nach Anspruch 1 oder 2, worin das Aminsalz Triethylaminhydrochlorid ist.
4. Verfahren nach irgendeinem der vorhergehenden Ansprüche, worin das aromatische Kohlenwasserstoff-Lösungsmittel Toluol oder Xylol ist.
5. Verfahren zur Herstellung eines 1-(Tetrazolylbiphenylmethyl)imidazol-Derivats der Formel (5):



worin R² Wasserstoff oder eine C₁-C₄-Alkylgruppe ist, wobei das Verfahren umfaßt die Umsetzung von 1 mol 1-(Cyanobiphenylmethyl)imidazol-Derivat der Formel (4):



worin R¹ Wasserstoff oder eine C₁-C₄-Alkylgruppe ist, und 1 bis 5 mol eines anorganischen Azidsalzes der Formel (2):

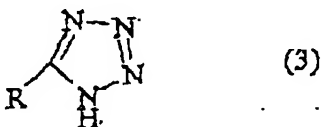


worin M ein Alkalimetall oder ein Erdalkalimetall ist und n 1 oder 2 ist, in einem aromatischen Kohlenwasserstoff-Lösungsmittel in Anwesenheit von 1 bis 5 mol eines Aminsalzes und gegebenenfalls das Hydrolysieren der erhaltenen Verbindung.

6. Verfahren nach Anspruch 5, worin R¹ in Formel (4) Methyl oder Ethyl ist und R² in Formel (5) Wasserstoff ist.
7. Verfahren nach Anspruch 5 oder 6, worin das anorganische Azidsalz Natriumazid ist.
8. Verfahren nach irgendeinem der Ansprüche 5 bis 7, worin das Aminsalz Triethylaminhydrochlorid ist.
9. Verfahren nach irgendeinem der Ansprüche 5 bis 8, worin das aromatische Kohlenwasserstoff-Lösungsmittel Toluol oder Xylol ist.

Revendications

1. Procédé pour produire un tétrazole 5-substitué de formule (3):



dans laquelle R est un groupe aliphatique, un groupe alicyclique, un groupe aromatique, un groupe hétérocyclique ou un groupe aliphatique hétérocyclique, chaque groupe étant non substitué ou substitué par un atome d'halogène ou un groupe hydroxy, cyano, nitro, alcoxy en C₁-C₈ ou amino, ou R est un groupe cycloalkyle en C₆-C₁₂ ou alkyle en C₁-C₈ ramifié ou non ramifié, chacun d'eux étant non substitué ou substitué par un groupe phényle, biphényle ou naphthyle, où les groupes phényle, biphényle et naphthyle peuvent être non substitué ou substitué par un atome d'halogène ou un groupe hydroxy, cyano, nitro, alcoxy en C₁-C₈ ou amino, procédé qui comprend la réaction, dans un solvant hydrocarboné aromatique et en présence de 1 à 5 moles d'un sel d'amine, de 1 mole d'un nitrile de formule (1):

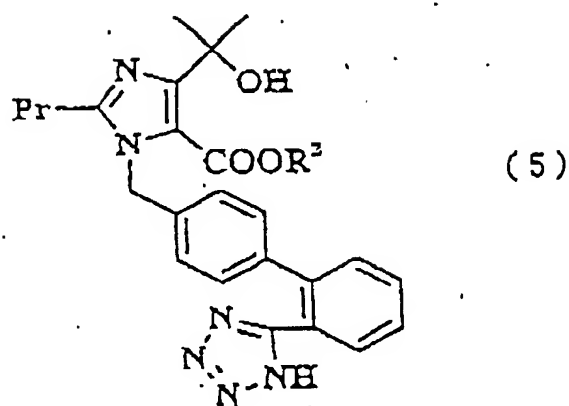


dans laquelle R est tel que défini ci-dessus, et de 1 à 5 moles d'un sel azoture minéral de formule (2):

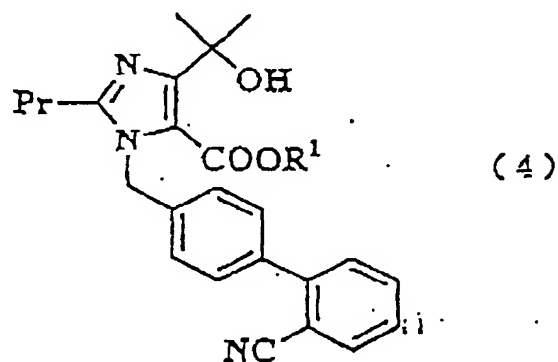


dans laquelle M est un métal alcalin ou un métal alcalino-terreux, et n vaut 1 ou 2.

2. Procédé selon la revendication 1, dans lequel le sel azoture minéral est l'azoture de sodium.
3. Procédé selon la revendication 1 ou 2, dans lequel le sel d'amine est le chlorhydrate de triéthylamine.
4. Procédé selon l'une quelconque des revendications précédentes, dans lequel le solvant hydrocarboné aromatique est le toluène ou le xylène.
5. Procédé pour produire un dérivé 1-(tétrazolylbiphénylméthyl)imidazole de formule (5):



20 dans laquelle R² est un atome d'hydrogène ou un groupe alkyle en C₁-C₄, procédé qui comprend la réaction, dans un solvant hydrocarboné aromatique et en présence de 1 à 5 moles d'un sel d'amine, de 1 mole d'un dérivé 1-(cyanobiphénylméthyl)imidazole de formule (4):



40 dans laquelle R¹ est un atome d'hydrogène ou un groupe alkyle en C₁-C₄, et de 1 à 5 moles d'un sel azoture minéral de formule (2):



dans laquelle M est un métal alcalin ou un métal alcalino-terreux, et n vaut 1 ou 2, et, le cas échéant, l'hydrolyse du composé obtenu.

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6. Procédé selon la revendication 5, dans lequel R¹, dans la formule (4), est un groupe méthyle ou éthyle, et R², dans la formule (5), est un atome d'hydrogène.
- 50
7. Procédé selon la revendication 5 ou 6, dans lequel le sel azoture minéral est l'azoture de sodium.
8. Procédé selon l'une quelconque des revendications 5 à 7, dans lequel le sel d'amine est le chlorhydrate de triéthylamine.
- 55
9. Procédé selon l'une quelconque des revendications 5 à 8, dans lequel le solvant hydrocarboné aromatique est le toluène ou le xylène.